SEARCH:

Title: US-10-590-810-26_COPY_13_555

Perfect score: 2763

Sequence: 1 SPKALEEAPWPPPEGAFVGF......LAVPLEVEVGIGEDWLSAKE 543

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 12150526 segs, 2531973831 residues

Total number of hits satisfying chosen parameters: 12150526

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100% Listing first 45 summaries

Database : Pending_Patents_AA_Main:*

SUMMARIES

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No.	Score		Length	DB	ID	Description
1	2763	100.0	543	27	US-09-791-537-91889	Sequence 91889, A
2	2763	100.0	544	39	US-10-917-157-2	Sequence 2, Appli
3	2763	100.0	544	39	US-10-917-157A-2	Sequence 2, Appli
4	2763	100.0	544	39	US-10-917-157B-2	Sequence 2, Appli
5	2763	100.0	545	39	US-10-917-157-4	Sequence 4, Appli
6	2763	100.0	545	39	US-10-917-157A-4	Sequence 4, Appli
7	2763	100.0	545	39	US-10-917-157B-4	Sequence 4, Appli
8	2763	100.0	552	25	US-09-506-153-1	Sequence 1, Appli
9	2763	100.0	552	32	US-10-216-682-1	Sequence 1, Appli
10	2763	100.0	554	1	PCT-US07-78571-3	Sequence 3, Appli
11	2763	100.0	554	12	US-08-202-032A-6	Sequence 6, Appli
12	2763	100.0	554	19	US-08-931-818-6	Sequence 6, Appli
13	2763	100.0	554	38	US-10-850-816-2	Sequence 2, Appli
14	2763	100.0	554	52	US-12-254-969B-6	Sequence 6, Appli
15	2763	100.0	554	53	US-12-330-201A-2	Sequence 2, Appli
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21	2763	100.0	562	32	US-10-216-682-7	Sequence 7, Appli
22	2763	100.0	562	35	US-10-590-810-26	Sequence 26, Appl
23	2763	100.0	605	43	US-11-327-195-44	Sequence 44, Appl
24	2763	100.0	605	43	US-11-327-195A-44	Sequence 44, Appl
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    44
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                                                           Sequence 8, Appli
                                    ALIGNMENTS
RESULT 1
US-09-791-537-91889
; Sequence 91889, Application US/09791537
; GENERAL INFORMATION:
; APPLICANT: Bionomix, Inc.
; APPLICANT: Debe, Derek
; APPLICANT: Danzer, Joseph
; TITLE OF INVENTION: THREE DIMENSIONAL STRUCTURES OF PROTEIN FAMILIES AND FAMILY
MEMBERS AND
; TITLE OF INVENTION: METHODS OF USE THEREOF
; FILE REFERENCE: 261/210
; CURRENT APPLICATION NUMBER: US/09/791,537
; CURRENT FILING DATE: 2001-02-22
; NUMBER OF SEO ID NOS: 153055
: SOFTWARE: PatentIn version 3.0
; SEO ID NO 91889
   LENGTH: 543
    TYPE: PRT
    ORGANISM: pdb 1KTQ
US-09-791-537-91889
  Query Match
                          100.0%; Score 2763; DB 27; Length 543;
  Best Local Similarity 100.0%;
  Matches 543; Conservative 0; Mismatches 0; Indels 0; Gaps
                                                                             0:
            1 SPKALEEAPWPPPEGAFVGFVLSRKEPMWADLLALAAARGGRVHRAPEPYKALRDLKEAR 60
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           61 GLIAKDLSVLALREGLGLPPGDDPMLLAYLLDPSNTTPEGVARRYGGEWTEEAGERAALS 120
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61 GLIAKDLSVLALREGLGLPPGDDPMLLAYLLDPSNTTPEGVARRYGGEWTEEAGERAALS 120

Qv

Db

Qv.

Db

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Db
Qy
          181 LEAEVFRLAGHPFNLNSRDQLERVLFDELGLPAIGKTEKTGKRSTSAAVLEALREAHPIV 240
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Db
          241 EKILOYRELTKLKSTYIDPLPDLIHPRTGRLHTRFNOTATATGRLSSSDPNLONIPVRTP 300
Qy
Dh
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          301 LGORIRRAFIAEEGWLLVALDYSOIELRVLAHLSGDENLIRVFOEGRDIHTETASWMFGV 360
Qу
Db
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Qy
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Db
          421 TLEEGRRRGYVETLFGRRRYVPDLEARVKSVREAAERMAFNMPVQGTAADLMKLAMVKLF 480
Qv
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          541 AKE 543
Οv
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21
1688
DMA
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(3)..(1688)
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  Met Ala Ser Gly Gly Gly Gly Cys Gly Gly Gly Gly Ser Pro Lys
gcc ctg gag gag gcc ccc tgg ccc ccg ccg gaa ggg gcc ttc gtg ggc
                                                              95
Ala Leu Glu Glu Ala Pro Trp Pro Pro Pro Glu Gly Ala Phe Val Gly
              20
                                25
ttt gtg ctt tcc cgc aag gag ccc atg tgg gcc gat ctt ctg gcc ctg
Phe Val Leu Ser Arg Lys Glu Pro Met Trp Ala Asp Leu Leu Ala Leu
                            40
          35
gcc gcc gcc agg ggg ggc cgg gtc cac cgg gcc ccc gag cct tat aaa
                                                             191
Ala Ala Ala Arg Gly Gly Arg Val His Arg Ala Pro Glu Pro Tyr Lys
                        55
gcc ctc agg gac ctg aag gag gcg cgg ggg ctt ctc gcc aaa gac ctg
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Ala Leu Arg Asp Leu Lys Glu Ala Arg Gly Leu Leu Ala Lys Asp Leu
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age gtt etg gee etg agg gaa gge ett gge ete eeg eee gge gae gae
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						ctc Leu										335
				100					105					110		
						ggc										383
Gly	Val	Ala	Arg 115	Arg	Tyr	Gly	Gly	Glu 120	Trp	Thr	Glu	Glu	Ala 125	Gly	Glu	
cgg	gcc	gcc	ctt	tcc	gag	agg	ctc	ttc	gcc	aac	ctg	tgg	ggg	agg	ctt	431
Arg	Ala	Ala 130	Leu	Ser	Glu	Arg	Leu 135	Phe	Ala	Asn	Leu	Trp 140	Gly	Arg	Leu	
gag	ggg	gag	gag	agg	ctc	ctt	tgg	ctt	tac	cgg	gag	gtg	gag	agg	ccc	479
Glu	Gly 145	Glu	Glu	Arg	Leu	Leu 150	Trp	Leu	Tyr	Arg	Glu 155	Val	Glu	Arg	Pro	
						cac										527
Leu 160	Ser	Ala	Val	Leu	Ala 165	His	Met	Glu	Ala	Thr 170	Gly	Val	Arg	Leu	Asp 175	
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Val	Ala	Tyr	Leu	Arg 180	Ala	Leu	Ser	Leu	Glu 185	Val	Ala	Glu	Glu	Ile 190	Ala	
cgc	ctc	gag	gcc	gag	gtc	ttc	cgc	ctg	gcc	ggc	cac	ccc	ttc	aac	ctc	623
Arg	Leu	Glu	Ala 195	Glu	Val	Phe	Arg	Leu 200	Ala	Gly	His	Pro	Phe 205	Asn	Leu	
aac	tcc	cgg	gac	cag	ctg	gaa	agg	gtc	ctc	ttt	gac	gag	cta	ggg	ctt	671
Asn	Ser	Arg 210	Asp	Gln	Leu	Glu	Arg 215	Val	Leu	Phe	Asp	Glu 220	Leu	Gly	Leu	
ccc	gcc	atc	ggc	aag	acg	gag	aag	acc	ggc	aag	cgc	tcc	acc	agc	gcc	719
Pro	Ala 225	Ile	Gly	Lys	Thr	Glu 230	Lys	Thr	Gly	Lys	Arg 235	Ser	Thr	Ser	Ala	
gcc	gtc	ctg	ggg	gcc	ctc	cgc	gag	gcc	cac	ccc	atc	gtg	gag	aag	atc	767
Ala 240	Val	Leu	Gly	Ala	Leu 245	Arg	Glu	Ala	His	Pro 250	Ile	Val	Glu	Lys	11e 255	
ctg	cag	tac	cgg	gag	ctc	acc	aag	ctg	aag	agc	acc	tac	att	gac	ccc	815
Leu	Gln	Tyr	Arg	Glu 260	Leu	Thr	Lys	Leu	Lys 265	Ser	Thr	Tyr	Ile	Asp 270	Pro	
ttg	ccg	gac	ctc	atc	cac	ccc	agg	acg	ggc	cgc	ctc	cac	acc	cgc	ttc	863
			275			Pro		280					285			
aac	cag	acg	gcc	acg	gcc	acg	ggc	agg	cta	agt	agc	tcc	gat	ccc	aac	911
Asn	Gln	Thr 290	Ala	Thr	Ala	Thr	Gly 295	Arg	Leu	Ser	Ser	Ser 300	Asp	Pro	Asn	
						cgc										959
	305					Arg 310					315					
						ggg										1007
	Phe	Ile	Ala	Glu		Gly	Trp	Leu	Leu		Thr	Leu	Asp	Tyr		
320					325					330					335	
						ctg										1055
				340		Leu			345		_	-		350		
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			355			Gly		360					365			
						cgg										1151
		370				Arg	375					380				
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Ala	Ala 385	Lys	Thr	Ile	Asn	Phe 390	Gly	Val	Leu	Tyr	Gly 395	Met	Ser	Ala	His	
cgc	ctc	tcc	cag	gag	cta	gcc	atc	cct	tac	gag	gag	gcc	cag	gcc	ttc	1247
	Leu	Ser	Gln	Glu		Ala	Ile	Pro	Tyr		Glu	Ala	Gln	Ala		
400					405					410					415	
att	gag	cgc	tac	ttt	cag	agc	ttc	ccc	aag	gtg	cgg	gcc	tgg	att	gag	1295

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aag acc ctg gag gag ggc agg agg cgg ggg tac gtg gag acc ctc ttc
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Lys Thr Leu Glu Glu Gly Arg Arg Gly Tyr Val Glu Thr Leu Phe
           435
                               440
                                                   445
ggc cgc cgc cgc tac gtg cca gac cta gag gcc cgg gtg aag agc gtg
                                                                    1391
Gly Arg Arg Arg Tyr Val Pro Asp Leu Glu Ala Arg Val Lys Ser Val
                           455
                                               460
cgg gag gcg gcc gag cgc atg gcc ttc aac atg ccc gtc cag ggc acc
                                                                    1439
Arg Glu Ala Ala Glu Arg Met Ala Phe Asn Met Pro Val Gln Gly Thr
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                                           475
gcc gcc gac ctc atg aag ctg gct atg gtg aag ctc ttc ccc agg ctg
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Ala Ala Asp Leu Met Lys Leu Ala Met Val Lys Leu Phe Pro Arg Leu
                   485
                                       490
gag gaa atg ggg gcc agg atg ctc ctt cag gtc cac gac gag ctg gtc
                                                                    1535
Glu Glu Met Gly Ala Arg Met Leu Leu Gln Val His Asp Glu Leu Val
                500
                                   505
ctc gag gcc cca aaa gag ggg gcg gag gcc gtg gcc cgg ctg gcc aag
                                                                    1583
Leu Glu Ala Pro Lys Glu Gly Ala Glu Ala Val Ala Arg Leu Ala Lys
           515
                               520
gag gtc atg gag ggg gtg tat ccc ctg gcc gtg ccc ctg gag gtg gag
                                                                    1631
Glu Val Met Glu Gly Val Tyr Pro Leu Ala Val Pro Leu Glu Val Glu
        530
                            535
gtg ggg ata ggg gag gac agg ctc tcc gcc aag gag gcg gcc gca ctg
                                                                    1679
Val Gly Ile Gly Glu Asp Arg Leu Ser Ala Lys Glu Ala Ala Ala Leu
   545
                       550
                                           555
                                                                    1688
gtg ccg cgc
Val Pro Arg
560
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W550 tryptophan

SEQ ID NO:26 is Thermus aquaticus

EAST SEARCH:

US 7488816 11/065,943 (Wilder exmnr)

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                     ("7417133").PN. US-PGPUB: USPAT: USOCR 2010/07/12 10:22
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d his

(FILE 'HOME' ENTERED AT 11:32:56 ON 12 JUL 2010)

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FILE 'MEDLINE, CAPLUS, BIOSIS, BIOTECHNO, EMBASE, JAPIO' ENTERED AT 11:36:07 ON 12 JUL 2010
L1 1642496 S POLYMERASE
L2 0 S L1 AND W550
L3 4979 S L1 AND TRYPTOPHAN
L4 3 S L3 AND 550
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1.5
          1522 S L1 AND 550
L6
          2207 S L3 AND (MUTA? OR VARIAN? OR SUBSTIT?)
L7
             12 S L6 AND TAO
1.8
              8 DUP REM L7 (4 DUPLICATES REMOVED)
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L12
              2 S L10 AND TAO
L8 ANSWER 4 OF 8 CAPLUS COPYRIGHT 2010 ACS on STN
AN 2004:171780 CAPLUS
DN
   140:351362
ΤI
    A novel strategy to engineer DNA polymerases for enhanced processivity and
     improved performance in vitro
    Wang, Yan; Prosen, Dennis E.; Mei, Li; Sullivan, John C.; Finney, Michael;
AU
    Vander Horn, Peter B.
    Department of Research and Development, MJ Bioworks Inc., South San
    Francisco, CA, 94080, USA
    Nucleic Acids Research (2004), 32(3), 1197-1207
SO
    CODEN: NARHAD; ISSN: 0305-1048
PR
    Oxford University Press
DT
    Journal
LA
    English
AB
   Mechanisms that allow replicative DNA polymerases to attain high
    processivity are often specific to a given polymerase and cannot
    be generalized to others. Here the authors report a protein
    engineering-based approach to significantly improve the processivity of
     DNA polymerases by covalently linking the polymerase domain to a
    sequence non-specific dsDNA binding protein. Using Sso7d from Sulfolobus
    solfataricus as the DNA binding protein, the authors demonstrate that the
    processivity of both family A and family B polymerases can be
    significantly enhanced. By introducing point mutations in
     Sso7d, the authors show that the dsDNA binding property of Sso7d is
     essential for the enhancement. The authors present evidence supporting
    two novel conclusions. First, the fusion of a heterologous dsDNA binding
    protein to a polymerase can increase processivity without
    compromising catalytic activity and enzyme stability. Second,
    polymerase processivity is limiting for the efficiency of PCR,
    such that the fusion enzymes exhibit profound advantages over unmodified
    enzymes in PCR applications. This technol, has the potential to broadly
     improve the performance of nucleic acid modifying enzymes.
OSC.G 26
             THERE ARE 26 CAPLUS RECORDS THAT CITE THIS RECORD (27 CITINGS)
RE.CNT 53
             THERE ARE 53 CITED REFERENCES AVAILABLE FOR THIS RECORD
             ALL CITATIONS AVAILABLE IN THE RE FORMAT
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